## IN THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in the application:

- 1-12 (Canceled)
- 13. (Withdrawn) A method of specifically inhibiting Rsk activity, said method comprising the step of contacting a Rsk enzyme with a compound represented by the general structure:

HO OH 
$$R$$
  $R_1$   $R_2$   $R_3$ 

wherein R is H or OH, and  $R_1$ ,  $R_2$  and  $R_3$  are independently selected from the group consisting of hydroxy -OCOR<sub>4</sub>, -COR<sub>4</sub> and  $C_1$ -C<sub>4</sub> alkoxy; and

 $R_4$  is H or  $C_1$ - $C_4$  alkyl.

- 14. (Withdrawn) The method of claim 13, wherein R is H or OH and R<sub>1</sub>, R<sub>2</sub> and R<sub>3</sub> are independently selected from the group consisting of hydroxy and -OCOCH<sub>3</sub>.
  - 15. (Withdrawn) The method of claims 13 or 14 wherein R is H.
  - 16. (Withdrawn) The method of claims 13, 14 or 15, wherein R<sub>3</sub> is -OCOCH<sub>3</sub>.
  - 17-20. (Canceled).
- 21. (Currently Amended) A method for treating a cancer characterized by excessive Rsk activity, said method comprising the step of administering to a human or other mammal, in

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need thereof having a cancer characterized by excessive Rsk activity and selected from the group consisting of breast, prostate, leukemia, lung, colon, brain, melanoma, ovarian, and kidney, a composition comprising a compound represented by the general structure:

HO OH O R
$$R_1$$

$$R_2$$

$$R_3$$

wherein R is H or OH, and R<sub>1</sub>, R<sub>2</sub> and R<sub>3</sub> are independently selected from the group consisting of hydroxy, -OCOCH<sub>3</sub>, -COCH<sub>3</sub>, C<sub>1</sub>-C<sub>4</sub> alkoxy, -O-glucoside and -O-rhamnoside in an amount effective for specifically inhibiting Rsk activity in the cells of said human or mammal.

- 22. (Original) The method of claim 21 wherein R is H and R<sub>1</sub>, R<sub>2</sub> and R<sub>3</sub> are independently selected from the group consisting of hydroxy and -OCOCH<sub>3</sub>.
- 23. (Original) The method of claim 21 wherein R is H or OH, R<sub>1</sub> and R<sub>2</sub> are independently hydroxy or -OCOCH3 and R3 is -OCOCH3.
  - 24. (Original) The method of claim 23 wherein R is H.
- 25. (Currently Amended) A method for treating a cancer characterized by excessive Rsk activity, said method comprising the step of administering to a patient, in need thereof having a cancer characterized by excessive Rsk activity and selected from the group consisting of breast, prostate, leukemia, lung, colon, brain, melanoma, ovarian, and kidney, a composition comprising a Rsk specific inhibitor in an amount effective for specifically inhibiting Rsk activity.

- 26. (Original) The method of claim 25 wherein the Rsk specific inhibitor comprises a compound selected from the group consisting of an anti-sense oligonucleotide and an interfering oligonucleotide.
- 27. (Original) The method of claim 25 wherein the Rsk specific inhibitor comprises an interfering oligonucleotide directed against Rsk1, Rsk2, Rsk3 or Rsk4.
- 28. (Original) The method of claim 25 wherein the Rsk specific inhibitor comprises an extract from the tissues of *Forsteronia refracta* or *Zingiber zerumbet*.
  - 29-31. (Canceled)
- 32. (Currently Amended) The method of claim 21 25 further comprising the steps of administering said human or other mammal patient an additional anti-tumor therapy.
  - 33-50 (Canceled)
- 51. (Previously presented) The method of claim 21, wherein said cancer is selected from the group consisting of breast cancer, prostate cancer, and sarcoma.
- 52. (Previously presented) The method of claim 51, wherein said Rsk inhibitor is selected from the group consisting of SL0101-1, SL0101-2, and SL0101-3, having the following structures:

and

SL0101-1

SL0101-3

53. (Previously presented) The method of claim 25, wherein said cancer is selected from the group consisting of breast cancer, prostate cancer, and sarcoma.

SL0101-2